Notice of References Cited Application/Control No. O9/503,758 Page 1 of 3 Applicant(s)/Patent Under Reexamination THILLY, WILLIAM G. Art Unit Page 1 of 3

U.S. PATENT DOCUMENTS

*		Document Number Country Code-Number-Kind Code	Date MM-YYYY	Name	Classification
	Α	US-			
	В	US-			
	С	US-			
	D	US-			
	Ε	US-			
	F	US-			
	G	US-			
	н	US-			
	1	US-			
	J	US-			
	К	US-			·
	L	US-			
	М	US-			

FOREIGN PATENT DOCUMENTS

*		Document Number Country Code-Number-Kind Code	Date MM-YYYY	Country	Name	Classification
	N					
	0					
	Р					
	Q					
<u> </u>	R					
	s					
	Т					

NON-PATENT DOCUMENTS

_		
*		Include as applicable: Author, Title Date, Publisher, Edition or Volume, Pertinent Pages)
	U	Davies, K. E. et al., "Molecular Basis of Inherited Disease", IRL Press, New York, pp, 21-25 (1992).
	V	Crow, J.F. et al., "Mutation in Human Population", Adv. Hum. Genet., vol. 14, pp. 59-77 (1985).
	w	Conneally, P. M., "Human Genetic Polymorphisms", Dev. Biol. Stand., vol. 83, pp. 107-110 (1994).
	x	de la Chapelle, A., "Disease gene mapping in isolated human populations; the example of Finland", J. Med. Genet., vol. 30, pp 857-865 (1993).

*A copy of this reference is not being furnished with this Office action. (See MPEP § 707.05(a).)

Dates in MM-YYYYY format are publication dates. Classifications may be US or foreign.

Notice of References Cited

Application/Control No.

O9/503,758

Applicant(s)/Patent Under Reexamination THILLY, WILLIAM G.

Examiner

Teresa E Strzelecka

Art Unit
Page 2 of 3

U.S. PATENT DOCUMENTS

*		Document Number Country Code-Number-Kind Code	Date MM-YYYY	Name	Classification
	A	US-			
	В	US-			
	С	US-		,	
,	D	US-			
	Е	US-			
	F	US-			·
	G	US-			·
	Н	US-			
	-	US-			
	j	US-			
	К	US-			
	L	US-			
	М	US-			

FOREIGN PATENT DOCUMENTS

*		Document Number Country Code-Number-Kind Code	Date MM-YYYY	Country	Name	Classification
	Ν					
	0					
	ъ					
	α					
	R		•			
	S					
	Т					

NON-PATENT DOCUMENTS

		NON-ATENT BOODINENTS
*		Include as applicable: Author, Title Date, Publisher, Edition or Volume, Pertinent Pages)
	U	Cavalli-Sforza, L. L. et al. "The Genetics of Human Populations", W. H. Freeman and Company, San Francisco, pp. 71-110 (1971).
	٧	Hardelin, J-P. et al., "Heterogeneity in the mutations responsible for X-chromosome linked Kallman syndrome", Hum. Mol. Genetics, vol. 2, pp. 373-377 (1993).
	w	Cooper, D. N. et al., "The mutational spectrum of single base-pair substitutions causing human genetic disease: patterns and predictions", Hum. Genetics, vol. 85, pp. 55-74 (1990).
	х	Maraglione, M. et al., "Prevalence of Apolipoprotein E Alleles in Healthy Subjects and Survivors of Ischemic Stroke", Stroke, vo 29, pp. 399-403 (February 1998).

*A copy of this reference is not being furnished with this Office action. (See MPEP § 707.05(a).)

Dates in MM-YYYY format are publication dates. Classifications may be US or foreign.